PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION See Form PCT/IPEA/416			
W 3435-004	International filing date (day/month/year)	Priority date (day/month/year)		
International application No.		17-02-2004		
	CT/SE2005/000220 17-02-2005 17-02-2004 remational Patent Classification (IPC) or national classification and IPC			
	or national classification and IPC			
See Supplemental Box				
Applicant				
Synbiotics AB et al				
This report is the international preliminary examination report, established by this International Preliminary Examining Authority under Article 35 and transmitted to the applicant according to Article 36.				
2. This REPORT consists of a total of				
This report is also accompanied b				
	t and to the International Bureau) a total of			
sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions).				
		rity considers contain an amendment that goes		
beyond the disclosure in the international application as filed, as indicated in item 4 of Box No. I and the Supplemental Box.				
I — — ¨		•		
b. (sent to the Internation	onal Bureau only) a total of (indicate type and			
, containing a sequence listing and/or tables related thereto, in electronic form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).				
This report contains indications re	elating to the following items:			
•				
Box No. II Priority	y			
	stablishment of opinion with regard to novelty,	inventive step and industrial applicability		
Box No. IV Lack of	f unity of invention			
Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial				
applicability; citations and explanations supporting such statement				
	Box No. VI Certain documents cited Box No. VII Certain defects in the international application			
Box No. VIII Certain observations on the international application				
Date of submission of the demand	Date of submission of the demand Date of completion of this report			
09.09.2005 24-02-2006				
Name and mailing address of the IPEA/S	SE Authorized officer	•		
Patent- och registreringsverket Box 5055				
S-102 42 STOCKHOLM	•	Terese Sandström/EÖ		
Facsimile No. +46 8 667 72 88	Telephone No. +4	Telephone No. +46 8 782 25 00		

Form PCT/IPEA/409 (cover sheet) (April 2005)

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Supplemental Box

In case the space in any of the preceding	boxes is not sufficient.

Continuation of: Cover sheet

INTERNATIONAL PATENT CLASSIFICATION (IPC):

A61K 35/74 (2006.01) A61P 29/00 (2006.01)

International application No.

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Box	No. I	Basis of the report					
1.	With re	egard to the language, this report is based on:					
	the international application in the language in which it was filed						
	a translation of the international application into						
	which is the language of a translation furnished for the purposes of:						
		publication of the international application (Rule	international search (Rules 12.3(a) and 23.1(b))				
		international preliminary examination (Rules 55.					
2.	furnish	regard to the elements of the international application, hed to the receiving Office in response to an invitation under not annexed to this report):	this report is based on (replacement sheets which have been ader Article 14 are referred to in this report as "originally filed"				
		the international application as originally filed/furnished					
1	\boxtimes	the description:					
		pages <u>1-18</u>	as originally filed/furnished				
		pages*rece	ived by this Authority onived by this Authority on				
			aveca by this Authority on				
	\bowtie	the claims:	as originally filed/furnished				
		pages*	as amended (together with any statement) under Article 19				
			eived by this Authority on 20-02-2006				
			eived by this Authority on				
		the drawings:					
		pages					
1		pages* rece	eived by this Authority on				
		•	eived by this Authority on				
!	Ш	a sequence listing and/or any related table(s) - see Supp	lemental Box Relating to Sequence Listing.				
3.		The amendments have resulted in the cancellation of:					
		the description, pages					
		the claims, Nos.					
		the drawings, sheets/figs					
Ì		the sequence listing (specify):					
ł		any table(s) related to the sequence listing (sp	ecify):				
4.		This report has been established as if (some of) the a made, since they have been considered to go beyond t 70.2(c)).	mendments annexed to this report and listed below had not been the disclosure as filed, as indicated in the Supplemental Box (Rule				
		the description, pages					
Ī		the claims, Nos.					
		the drawings, sheets/figs					
		the sequence listing (specify):					
		any table(s) related to the sequence listing (sp					
*	If iter	m 4 applies, some or all of those sheets may be marked "s	uperseded."				

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INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

	PC1/5E2005/000220
Box No. II Priority	
1. This report has been established as if no priority had been claimed due to the f limit the requested:	ailure to furnish within the prescribed time
copy of the earlier application whose priority has been claimed (Rule 66	.7(a)).
translation of the earlier application whose priority has been claimed (Ru	ule 66.7(b)).
2. This report has been established as if no priority had been claimed due to the finvalid (Rule 64.1). Thus for the purposes of this report, the international filing relevant date.	fact that the priority claim has been found g date indicated above is considered to be the
3. Additional observations, if necessary:	
matter considering a composition of two specified lactic acid bacteria the claim valid. However, since the present claims as	a for preventing or orders whereas the at least two of the venting or treating. Thus, for subject of or three of the ned priority is not all restricted to sified lactic acid biotic Modulation of Encephalopathy in wril 2004, Vol. 39, Changes in cytokine utrition with lactic or major abdominal .8, Suppl. 1, P273' tions comprising the

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Box No.	III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
The quest applicable	tions whether the claimed invention appears to be novel, to involve an inventive step (to be non obvious), or to be industrially e have not been examined in respect of:
	the entire international application
\boxtimes	claims Nos. 12
becau	se:
ani	the said international application, or the said claims Nos. 12 relate to the following subject matter which does not require an international preliminary examination (specify): PCT Rule 67.1.(iv).: Methods for treatment of the human or mal body by surgery or therapy, as well as diagnostic hods.
	the description, claims or drawings (indicate particular elements below) or said claims Nosare so unclear that no meaningful opinion could be formed (specify):
	the claims, or said claims Nos are so inadequately supported by the description that no meaningful opinion could be formed (specify):
	no international search report has been established for said claims Nos.
	a meaningful opinion could not be formed without the sequence listing; the applicant did not, within the prescribed time limit:
	furnish a sequence listing on paper complying with the standard provided for in Annex C of the Administrative Instructions, and such listing was not available to the International Preliminary Examining Authority in a form and manner acceptable to it. furnish a sequence listing in electronic form complying with the standard provided for in Annex C of the Administrative Instructions, and such listing was not available to the International Preliminary Examining Authority in a form and manner acceptable to it. pay the required late furnishing fee for the furnishing of a sequence listing in response to an invitation under Rules 13ter.1(a) or (b) and 13ter.2.
	a meaningful opinion could not be formed without the tables related to the sequence listings; the applicant did not, within the prescribed time limit, furnish such tables in electronic form complying with the technical requirements provided for in Annex C-bis of the Administrative Instructions, and such tables were not available to the International Preliminary Examining Authority in a form and manner acceptable to it.
	the tables related to the nucleotide and/or amino acid sequence listing, if in electronic form only, do not comply with the technical requirements provided for in the Annex C-bis of the Administrative Instructions.
	See Supplemental Box for further details.

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Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
1. Statemen	t			
Nove	elty (N)	Claims Claims	1-11	YES NO
Inver	ntive step (IS)	Claims Claims	1-11	YES NO
Indu	strial applicability (IA)	Claims Claims	1-11	YES NO

2. Citations and explanations (Rule 70.7)

The present claims relate to the use of a composition comprising the four lactic acid bacteria Pediococcus pentosecus 5-33:3, Leuconostoc mesenteroides 32-77:1, L. paracasei subsp. paracasei 19 and L. plantarum 2362, wherein the bacterial strains are in an amount of at least 10¹¹ CFU/ml of each of the bacteria, in combination with at least four different fibres for the manufacturing of a formulation for the prevention of a stress-induced inflammatory disorder.

Documents cited in the International Search Report:

- D1: Colucci G. et al., "Prevention of Postoperative Adhesions after Abdominal Aortic Surgery", Eur. Surg. Res., 2003, Vol. 35, page 265; P25
- D2: Bengmark S., "Use of some pre-, pro- and symbiotics in critically ill patients", Best Practice & Research Clinical Gastroenterology, 2003, Vol. 17, No. 5, pages 833-848
- D3: Bengmark S. "Modulation by enteral nutrition of the acute phase response and immune functions", Nutr. Hosp., 2003, Vol. 18, No. 1, pages 1-5
- D4: Bengmark S., "Symbiotic Control of Inflammation and Infection in Transplantation", Transplantation Reviews, January 2004, Vol. 18, No. 1, pages 38-53
- D1 discloses the administration of "Symbiotic 2000" to mice undergoing surgery. It was demonstrated that the use of "Symbiotic 2000" reduced inflammation response in those mice. "Symbiotic 2000" consists of the four lactic acid bacteria Pediococcus pentosecus 5-33:3, Leuconostoc mesenteroides 32-77:1, L. paracasei subsp. paracasei 19 and L. plantarum 2362 in combination with the four fermentable fibres beta-glucan, inulin, pectin and resistant starch (see e.g. D2 for the specific fibres). The composition in D1 includes 108 CFU of each of the lactic acid bacteria.

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Supplemental Box

In case the space in any of the preceding boxes is not sufficient. Continuation of: Box V

D2 discloses the use of "Symbiotic 2000" for the treatment of chronic distal colitis. Patients with chronic distal colitis treated with "Symbiotic 2000", administered twice-daily enemas, showed e.g. significant reductions in diarrhoea scores as well as visible blood in the stools. (Page 844, paragraph 2.) The composition used included 10¹⁰ CFU of each of the four lactic acid bacteria.

D3 suggests the use of "Synbiotic 2000" to modulate the acute phase response and limit induced superinflammation. (Page 4, column 1, last paragraph -column 2, paragraph 1.) D3 does not disclose the amount of bacterial strains in the composition. However, looking at the state of the art, "Synbiotic 2000" seems to refer to a composition comprising the 4 bacterial strains in amounts of 10⁸ or 10¹⁰ CFU/ml.

D4 suggests the use of symbiotic in general for reducing liver. D4 discloses some symbiotic inflammation of the composition comprising 1 lactic one compositions, bacteria in combination with one fibre and two compositions comprising 4 lactic acid bacteria in combination with four fibres ("Synbiotic 2000" and "Synbiotic FORTE"). "Synbiotic FORTE" differs from "Symbiotic 2000" since it comprises 1011 of each of the four lactic acid bacteria. The composition comprising one lactic acid bacteria in combination with one fibre was shown to treat pancreatitis as well as to reduce inflammation in the liver. (Page 48, column 1, paragraph 1; page 49, column 1, last 4 lines.)

The use as claimed in the present claims is not disclosed by any of the documents D1-D4. None of the documents describes the use of the four specific bacterial strains as defined in the claims in an amount of at least 10¹¹ CFU/ml of each of the bacteria and at least four fibres for the manufacture of a formulation for the prevention, i.e. not the treatment, of a stress-induced inflammatory disorder. Hence, the subject matter claimed in claims 1-11 is novel.

None of these documents are focused on solving the problem of preventing stress-induced inflammatory disorders, even though the inflammation in D1 might be considered to be a stress induced inflammatory disorder.

D4, which disclosed the composition "Symbiotic FORTE", is considered to be one document disclosing the closest prior art.

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In case the space in any of the preceding boxes is not sufficient. Continuation of: Box V

Since "Symbiotic FORTE", a composition comprising 1011 CFU/ml of each of the lactic acid bacteria, is one of the symbiotic compositions reviewed in the article, it might seem close to hand for a person skilled in the art, wishing to reduce inflammations in general or even stress-induced inflammatory disorders, to use this symbiotic composition. However, person skilled in the art faced with the problem of preventing a stress-induced inflammatory disorder coming across D4 would assume that to prevent a disease instead of treating/curing a disease could easily be done by the use of solely combination of oat and L. plantarum (the composition used in D4 comprising one lactic acid bacteria and one fibre) and would not try to prepare a mixture of four different specific bacterial strains and four fibres in the large amount of 1011 CFU/ml (as in the case on the "synbiotic FORTE" composition). It is less expensive and less complicated to use a less complicated composition.

The main difference between the subject matter claimed in claims 1-11 and the prior art stated in D1-D4 is the high amount of the four bacterial strains.

According to the response of the applicant, the inventors have surprisingly found that the use of such high amounts of the four bacterial strains in combination with four fibres gives rise to the following activities within the person to which the compositions is administrated, making it suitable for the claimed use:

The expression of the heat shock proteins increases
The expression of the nuclear factor (NF) kappa B decreases
TNF-alfa, Interleukin (IL)-6 and other markers of inflammation
decreases

Caspase-3 cell and tissue expression decreases Human leukocyte antigen (HLA)-DR expression improves

The NF-kB expression is reduced

COX 2 expression is reduced

The iNOS expression is reduced

The PAI-1 activity if reduced

The tissue infiltration of neutrophils is prevented

Tissue destruction is prevented

Gut for a is restored

This difference is not predictable to a person skilled in the art in view of the documents cited above.

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Hence it is not considered obvious for a person skilled in the art to use the four specific bacterial strains as defined in the claims in an amount of at least 10^{11} CFU/ml of each of the bacteria and at least four fibres for the manufacture of a formulation for the prevention, i.e. not the treatment, of a stress-induced inflammatory disorder.

To summaries, the subject matter claimed in claims 1-11 is novel and is considered to involve an inventive step. The subject matter claimed in claims 1-11 is considered to be industrially applicable.

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1 xc	No. VI Certain document	ts cited			
	Certain published documents	(Rule 70.10)			
	Application No. Patent No.	Pu <i>(dd</i>	ablication date sy/month/year)	Filing date (day/month/year)	Priority date (valid claim) (day/month/year)
	WO2004103083	A1 12	2.12.2004	18.05.2004	22.05.2003
_	Non-written disclosures (Rule	e 70.9)			Date of written disclosure
	Kind of non-writter	disclosure	Date of non- (day/n	written disclosure nonth/year)	referring to non-written disclosure (day/month/year)
	•				

CLAIMS

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- 1. Use of Pediococcus pentosaceus 16:1 (LMG P-20608), Leuconostoc mesenteroides 23-77:1 (LMG P-20607), Lactobacillus paracasei subsp paracasei F-19 (LMG P-17806), and Lactobacillus plantarum 2362 (LMG P-20606) wherein the bacterial strains are in an amount of at least 10¹¹ CFU/ml of each of the bacteria and at least four different fibres for the manufacturing of a formulation for the prevention of a stress-induced inflammatory disorder.
- 2. Use according to claim 1, wherein the stress-induced inflammatory disorder is determined as an increase in neutrophils, cytokines, myeloperoxidase and/or accumulation of the oxidation-related malonedealdehyde.
- 3. Use according to any of proceeding claims, wherein the inflammatory disorder is lung inflammation, urinary inflammation, vaginal inflammation, bowel inflammation, stomach inflammation, liver inflammation, muscle inflammation, inflammation of endocrine and reproductive organs, and brain inflammation.
- 4. Use according to claim 3, wherein the fibre is selected from the group consisting of beta-glucan, inulin, pectin, resistant starch, cellulose, hemicellulose, arabinoxylans, arabinogalactans, polyfructose, inulin, oligofructans, galacto-oligosacharides, gums, mucilages, pectins, dextrins, maltodextrins, potato dextrins, synthesised carbohydrates, polydextrose, methylcellulose and hydroxypropylmethylcellulose.
- 5. Use according to claim 4, wherein the four fibres are inulin, beta-glucan, pectin and resistant starch.
- 6. Use according to claim 5, wherein the fibres are present in an amount of 2.5 g of each fibre.
- 7. Use according to claim 4, wherein the fibre is selected from lignin substances from plants selected from the group comprising waxes, cutin, phytate, saponin, suberin and tannins.
- 8. Use according to any of proceeding claims, wherein the formulation further comprises at least one antioxidant, vitamin, mineral, amino acid, peptide or protein.
- 9. Use according to any of proceeding claims, wherein the formulation further comprises glutamine, or a synthetic version thereof.
- 10. Use according to any of proceeding claims, wherein the formulation further comprises one or more therapeutic agents.
- 11. Use according to any of preceding claims, wherein the formulation is solid or liquid, such as tablet, gel or spray.
- 12. Use of the formulation according to any of proceeding claims for the prevention of a mammal suffering from a stress-induced inflammatory disorder, such as an animal or human being.